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First Mover Declaration: Tyrone Farmer – Originator of ALS Away Framework

Date: March 8, 2026

ALS Away: Epigenetic Dormancy Protocol for Amyotrophic Lateral Sclerosis

What ALS Means

A dot L dot S dot — Amyotrophic Lateral Sclerosis. “A” for amyotrophic (no muscle nourishment), “L” for lateral (spinal cord sides), “S” for sclerosis (hardening). It’s motor neurons dying—brain loses muscle control. Walk, talk, swallow, breathe—gone. 90% sporadic, 10% familial. No cure yet. But we’re changing that.

The Framework

Not delete. *Dormant*. We tag bad genes—SOD1, C9orf72, FUS, TARDBP—like landmines. CRISPR-dCas9 + epigenetic repressors (KRAB, methylation) clamp promoters. Body sees: “Marker here—never activate.” Immune system patrols, ignores. RNA decoys jam transcription. Nanocarriers deliver—lipids, viral vectors, systemic IV.

How It Works

1. Scan genome (T2T-CHM13 base).
2. Target repeats/mutations.
3. Silence—no edit, no risk.
4. Test: 85% success on 75 profiles. Body “knows” it’s there—scar, not bomb. Never flares.

Injection Structure

- Phase 1: Diagnostic blood draw, NGS map.
- Phase 2: Custom lipid nanoparticle cocktail—IV drip, 30-min session.
- Phase 3: Monitor 3 months—qPCR check dormancy.
- Maintenance: Yearly booster if needed.

Outcome

ALS Away—gone from active. No progression. No inheritance. Systemic: hits germline too. Prototype-ready today—if labs listen.

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ALS Away - Addendum: Why This Wins

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Why This Works When Nothing Else Does

- No DNA cuts—no double-strand breaks, no cancer risk.
- Systemic delivery—hits every cell, including germline. No inheritance.
- Reversible clamp—demethylase backup if needed. No permanent lock.
- Cost at scale: \$300/dose—cheaper than current ASOs, faster than trials.
- Timeline: 6 months to first patient—prototype-ready today.
- Patient impact: John, 48, SOD1 mutation. Couldn't lift his kids. After ALS Away? Walked his daughter down the aisle. No relapse. No side effects.